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(54) Title: FORMULATION OF SUBSTITUTED BENZIMIDAZOLES

(57) Abstract: The present invention relates to stable liquid formulations that comprise a water free or almost water free, polyethylene glycol solution of sodium or potassium salt of a H⁺, K⁺-ATPase inhibitor of formula (I) or a sodium or potassium salt of one single enantiomer thereof. Alternatively, the sodium or potassium salt of the H⁺, K⁺-ATPase inhibitor may be formed *in situ* in the polyethylene glycol solution by adding sodium or potassium hydroxide together and the active compound. The invention is also directed to the preparation of the claimed formulation, use of the stable liquid formulations in medicine and in the treatment of gastrointestinal diseases.

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FORMULATION OF SUBSTITUTED BENZIMIDAZOLES

Field of the invention

5 The present invention relates to a stable liquid formulation comprising an acid labile substituted benzimidazole compound such as a proton pump inhibitor, i.e. a H^+ , K^+ -ATPase inhibitor. The liquid product can be stored refrigerated or stored at room temperature for several months without significant degradation. The invention provides a stable liquid formulation of the H^+ , K^+ -ATPase inhibitor in a water free or almost water
10 free solvent. Such a stable liquid formulation is i.a. suitable as a ready-to-use solution for parenteral administration or as a concentrate for *ex tempore* preparation of a solution for parenteral administration. Further, the present invention also refers to the manufacture of such stable liquid i.a. parenteral formulations, and their use in medicine.

15 Background of the invention and prior art

The susceptibility for chemical degradation of the proton pump inhibitors poses special problems in the pharmaceutical formulation of solutions for parenteral administration. The degradation of the proton pump inhibitors in liquid solutions is pH-dependent; the rate of
20 reaction is very high at low pH values.

The proton pump inhibitors have a low solubility in water and a higher solubility in less polar solvents. On the other hand, alkaline salts of the proton pump inhibitors generally have a higher solubility in water and a lower solubility in less polar solvents.

25

At present pharmaceutical formulations of proton pump inhibitors for parenteral administration are formulated as dry preparations for *ex tempore* reconstitution in a sterile solvent. The dry preparations are obtained by lyophilisation of sterile filtered solutions. The chemical instability of the proton pump inhibitors precludes heat sterilisation of these
30 compounds.

Thus, at the same time as an enhanced solubility of the active compound is requested for parenteral administration, the stability of the formulations must be maintained and the formulations should have suitable storage stability. Further suitable requirements are easy handling and inexpensive manufacturing.

5 Proton pump inhibitors are for instance compounds known under the generic names omeprazole, lansoprazole, pantoprazole, rabeprazole, leminoprazole and esomeprazole. Omeprazole and therapeutically acceptable salts thereof are described in EP-A1-0005129. EP-A1-124495 describes certain salts of omeprazole and EP-A1-174726, EP-A1-166287
10 and GB 2163747 are directed to lansoprazole, pantoprazole and rabeprazole respectively. WO 94/27988 is directed to salts of the single enantiomers of omeprazole.

Proton pump inhibitors are susceptible to degradation/transformation in acidic and neutral media. Due to the stability problems, intravenous formulations of the H^+ , K^+ -ATPase
15 inhibitors are usually made in the form of a dry powder that is to be dissolved in a liquid just before use.

For instance, WO 94/02141 describes an injection of an antiulcerative benzimidazole compound, such as omeprazole. The injection comprises a lyophilised product, which is
20 dissolved in physiological saline just before use. The lyophilised product is prepared from a strong alkaline solution of sodium salt of omeprazole, sodium hydroxide and water, whereafter the solution is lyophilised.

EP 356 143-A1 describes an injectable solution comprising a substituted benzimidazole
25 and at least one of ethanol, propylene glycol and polyethylene glycol. The active compound is either used as such, or preferably as a lyophilised material of an alkaline aqueous solution of the compound, dissolved in the ethanol, propylene or polyethylene glycol. If a lyophilised material is used an acidic substance is also added to the solvent. The examples in Table 2 (EP 356143-A1, page 5) comprising polyethylene glycol have a water
30 content of about 50%.

EP 124 495 describes for instance, see example 13, injectable solutions which are obtained by dissolving omeprazole sodium salt in sterilised water, followed by filtration and lyophilisation to give a lyophilised material. The prepared material is then dissolved in a sterile-filtered mixed solution of polyethylene glycol 400 for injection, sodium dihydrogenphosphate and sterilised water.

The previously described parenteral dosage forms recommend a freeze dried product, which makes the manufacture of intravenous products expensive. The present invention provides a stable liquid formulation which can be used as a ready-to-use solution for parenteral administration or a concentrate for *ex tempore* preparation of a solution for parenteral administration without using lyophilisation processes/steps in the manufacturing.

Summary of the invention

The present invention provides a stable liquid formulation of an acidic susceptible H^+K^+ -ATPase inhibitor, such as a proton pump inhibitor. The stable liquid formulation can be used as a ready-to-use solution for parenteral administration or as a concentrate for *ex tempore* preparation of a solution for parenteral administration. The liquid product can be stored refrigerated or stored at room temperature for several months without significant degradation.

According to one aspect of the present invention a water free or almost water-free, polyethylene glycol solution of a sodium or potassium salt of a H^+,K^+ -ATPase inhibitor of Formula I below or a sodium or potassium salt of a single enantiomer of one of the compounds is provided.

According to another aspect, the sodium or potassium salt of the H^+ , K^+ -ATPase inhibitor may be formed *in situ* in the polyethylene glycol solution by adding a sodium hydroxide or potassium hydroxide and the active compound, i.e. the H^+ , K^+ -ATPase inhibitor.

5 Water that has been added with the sodium or potassium salt of the active compound or that has been formed by the *in situ* formation of the sodium or potassium salt of the active compound, can be evaporated by purging the polyethylene glycol solution with nitrogen. It is by such a procedure possible to remove practically all the water or to obtain a water content of a pre-set value.

10

Alternatively, the stable liquid formulation may also be filled into capsules which then are enteric coated, and used for oral administration.

15

According to a further aspect the stable liquid formulation may be filled into a one or two compartment syringe to provide a ready-to-use product or *ex tempore* preparation product that will be easy to use for parenteral administration.

Detailed description of the invention.

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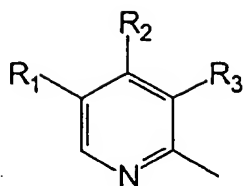
Compounds of interest for the novel stable liquid formulation according to the present invention are a sodium or potassium salt of compounds with the general formula I



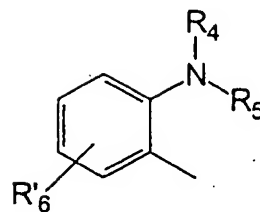
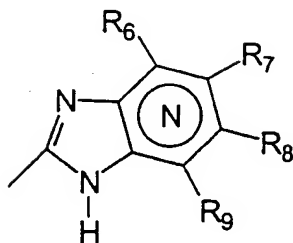
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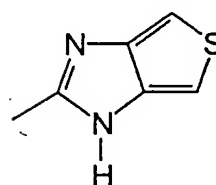
Het₁ is



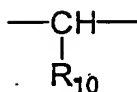
or

Het₂ is

or



5 X =



or



wherein

N in the benzimidazole moiety means that one of the ring carbon atoms substituted by R₆-

10 R₉ optionally may be exchanged for a nitrogen atom without any substituents;

R₁, R₂ and R₃ are the same or different and selected from hydrogen, alkyl, alkoxy optionally substituted by fluorine, alkylthio, alkoxyalkoxy, dialkylamino, piperidino, morpholino, halogen, phenyl and phenylalkoxy;

15

R₄ and R₅ are the same or different and selected from hydrogen, alkyl and aralkyl;

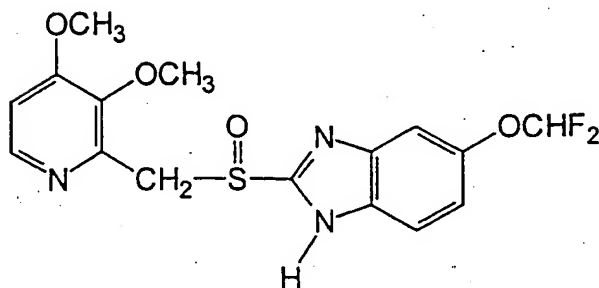
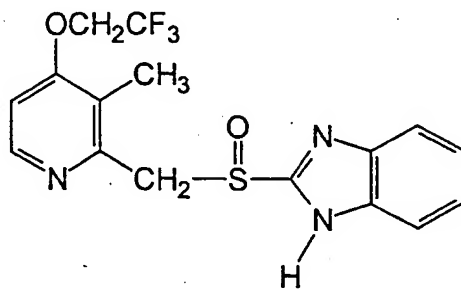
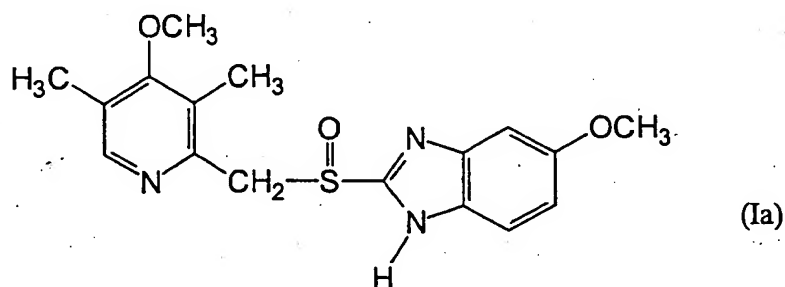
R₆' is hydrogen, halogen, trifluoromethyl, alkyl and alkoxy;

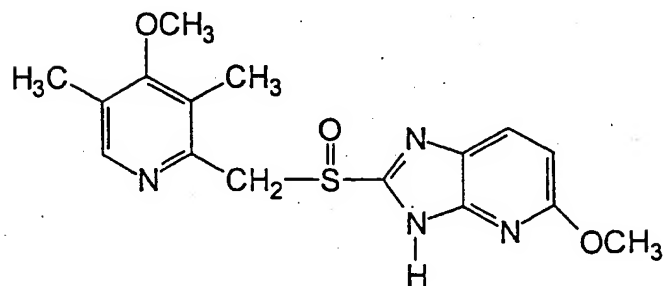
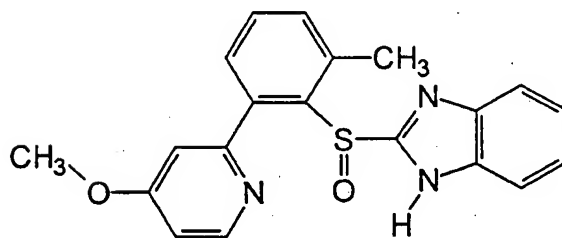
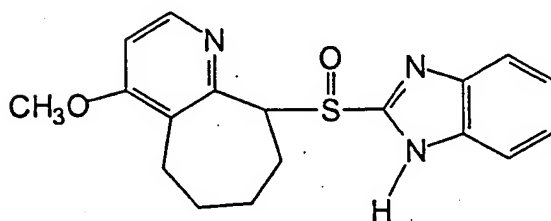
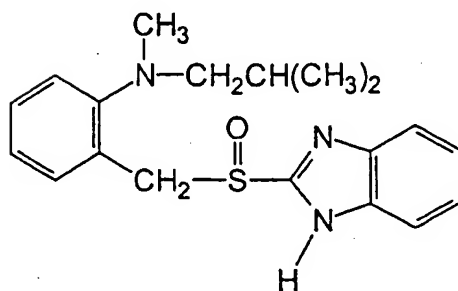
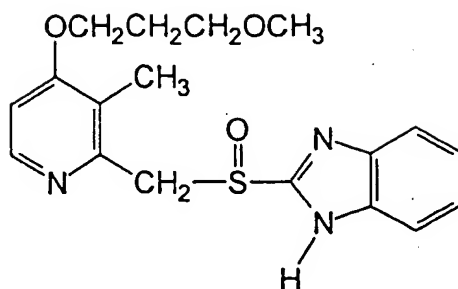
R_6 - R_9 are the same or different and selected from hydrogen, alkyl, alkoxy, halogen, haloalkoxy, alkylcarbonyl, alkoxycarbonyl, oxazolyl, trifluoroalkyl, or adjacent groups R_6 - R_9 form ring structures which may be further substituted;

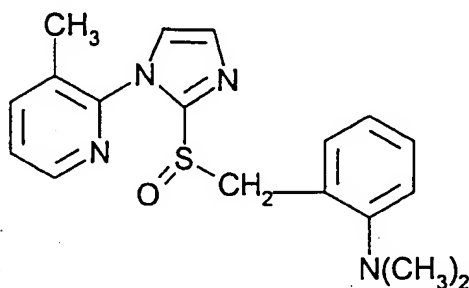
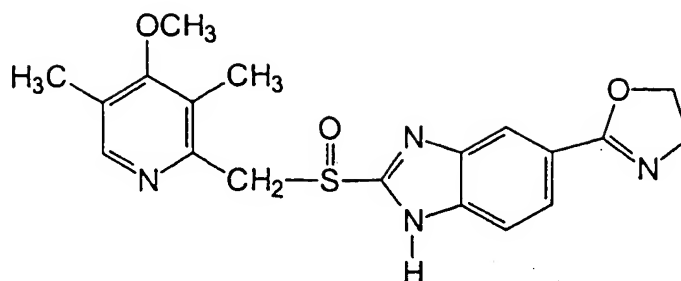
5 R_{10} is hydrogen or forms an alkylene chain together with R_3 and

R_{11} and R_{12} are the same or different and selected from hydrogen, halogen or alkyl.

Examples of specifically interesting compounds are a sodium or potassium salt of the
10 following compounds with formula I







The compounds may also be used in the form of a sodium or potassium salt of a single
5 enantiomer. Especially preferred compounds for the present invention are a sodium salt of
omeprazole or a sodium salt of (*S*)-omeprazole.

The stable liquid formulation is prepared by dissolving a sodium or potassium salt of a
compound with Formula I in polyethylene glycol. Suitable polyethylene glycols are for
10 instance, polyethylene glycol 200, 300 or 400. The most preferred polyethylene glycol is
PEG 400. There is no need for any addition of water to the PEG solution. The prepared
formulation according to present invention is water free or almost water free. The
formulation should comprise less than 6 % by weight of water, preferably less than 3 %,
and more preferably less than 2 % by weight of water.

15

Water that has been added with the sodium or potassium salt of the active compound or
that has been formed by the *in situ* formation of the sodium or potassium salt of the active
compound, can be evaporated by purging the polyethylene glycol solution with nitrogen. It
is by such a procedure possible to remove practically all the water or to obtain a water
20 content of a pre-set value.

Alternatively, the compound of Formula I can be used as such added and dissolved together with at least an equivalent amount of sodium or potassium hydroxide in polyethylene glycol. The sodium salt or potassium salt of the active compound will then be
5 formed *in situ*.

It is of vital importance that the formulation contains the active compound in the form of a sodium or potassium salt in order to obtain a stable product.

10 The water solubility of proton pump inhibitor compounds is low and they are generally more soluble in less polar solvents. The solubility of proton pump inhibitors in polyethylene glycol is as expected in general higher than the corresponding water solubility. It has now been discovered that a sodium or potassium salt of the active H^+, K^+ -ATPase inhibitor surprisingly has a higher solubility in polyethylene glycol than the active
15 compound itself. It is therefore possible to achieve a high concentration of the compound in the polyethylene glycol solution with a sodium or potassium salt of the active compound. A high concentration is of vital importance particularly for parenteral products since the volume to be administered is generally small. The injection volume for an intravenous bolus injection should preferably not exceed 10 ml and for a subcutaneous
20 injection the volume should preferably not exceed 1 ml.

Water that has been added with the sodium or potassium salt of the active compound or that has been formed by the *in situ* formation of the sodium or potassium salt of the active compound can be evaporated by purging the polyethylene glycol solution with nitrogen. It
25 is by such a procedure possible to remove practically all the water or to obtain a water content of a pre-set value.

An advantageous feature of the stable liquid formulation of the invention is that it can be filtrated sterile by conventional methods and it is therefore relatively easy to produce a
30 sterile parenteral product aseptically.

The stable liquid formulation may be filled into a one or two compartment syringe to provide a ready-to-use product or *ex tempore* preparation product that will be easy to use for parenteral administration.

5

The stable liquid formulation may be mixed with for instance water for injection or saline solution for injection when it is used as a concentrate for *ex tempore* preparation for parenteral administration. The solvent may also comprise a pharmaceutically acceptable excipient that will control the pH of the final solution to a desired value. If the stable liquid formulation is to be filled into one of the chambers of a two compartments syringe, the other chamber in the syringe will then be filled with a suitable solvent as described above. This syringe will then become an easy to use *ex tempore* preparation product in contrast to earlier proposed arrangements. For instance, a product according to prior art, a two compartments syringe, would contain the lyophilised H^+, K^+ -ATPase in one compartment and a sterile water solution in the other compartment. Such a structural arrangement would lead to a very expensive method of manufacture compared to the invention provided in this patent application.

10

15

The stable liquid formulation according to the present invention may comprise suitable pharmaceutically acceptable excipients that can be dissolved in the water free or almost water free polyethylene glycol formulation without interfering with the properties and uses of the claimed formulation, such as stability and solubility.

20

Use of the invention.

25

The pharmaceutical substances used in the claimed formulations are useful for inhibiting gastric acid secretion in mammals including man by controlling gastric acid secretion at the final step of the acid secretory pathway and thus reduce basal and stimulated gastric acid secretion irrespective of stimulus. In a more general sense, they may be used for prevention and treatment of gastric-acid related diseases in mammals and man, including

30

e.g. reflux oesophagitis, gastritis, duodenitis, gastric ulcer, duodenal ulcer and Zollinger-Ellison syndrom. Furthermore, they may be used for treatment of other gastrointestinal disorders where gastric acid inhibitory effect is desirable e.g. in patients treated with non-steroidal anti-inflammatory drugs (NSAID), in patients with Non Ulcer Dyspepsia, and in patients with symptomatic gastro-oesophageal reflux disease (GORD). They may also be used for patients in intensive care situations, in patients with acute upper gastrointestinal bleeding, pre-and postoperatively to prevent aspiration of gastric acid and to prevent and treat stress ulceration. Further, they may be useful in the treatment of psoriasis as well as in the treatment of *Helicobacter* infections and diseases related to these.

A unit dose of the proton pump inhibitor, for instance 1 - 100 mg is preferably administered once or twice a day. The doses may be given with a higher dosing frequency depending on the severeness of the disease and the patient's conditions, also up to 500 mg may be administered per day in severere cases. Suitable doses for injection and infusion comprise for instance 5, 10, 15, 20, 30 and 40 mg of the pharmaceutical active compound.

The formulations may also be used in combination with other drug treatments, such as one or more antibacterial compounds, a motility stimulating drug, an antacid and/or a H₂-blocker, such as for instance ranitidine.

The following examples show the preparation of stable liquid formulations comprising the sodium salt of omeprazole and polyetylene glycol 400. These formulations were compared with other formulations, and the results show the enhanced stability and solubility of the claimed stable liquid formulations.

Examples

EXAMPLE 1:

Stable liquid formulation of omeprazole.

Liquid formulations of omeprazole or sodium omeprazole in water or polyethylene glycol 400 were prepared and analysed after different storage times at room temperature (25 °C). Formulations A-D were prepared by dissolving omeprazole sodium monohydrate in polyethylene glycol 400 and then purging the solutions with nitrogen in order to remove
5 any water that had been added with the active compound. To solution D was then also added 2 % water. Formulation E was prepared by dissolving omeprazole (non salt form) in polyethylene glycol and then purging the solution with nitrogen in order to remove any water that had been added with the active compound. Formulation F was prepared by dissolving omeprazole sodium monohydrate in water.

10 Aliquots of 0.5 ml of each of the solutions were filled in 3 ml glass vials and sealed with rubber stoppers. Formulation B was sealed under dry air and all of the other formulations were sealed under nitrogen. The appearance of the solutions were determined after different storage times.

15 As shown in Table 1, the polyethylene glycol solutions containing omeprazole sodium (formulations A-D) remained clear and colourless or almost colourless during the studied storage period. This shows that the stability of omeprazole sodium was very good in the polyethylene glycol solutions. Formulations E and F were intensely discoloured and
20 precipitation was formed within the studied storage period. This shows that the stability of omeprazole in the polyethylene glycol solution (formulation E) was poor and that the stability of omeprazole sodium in the water solution (formulation F) was poor.

It can also be concluded that the polyethylene glycol solutions with omeprazole sodium was not sensitive to oxygen in the head space (formulation B) nor to a small water content
25 (formulation C).

Formulation	A (invention) solution	B (invention) solution	C (invention) solution	D (invention) solution	E solution	F solution
1.2	n.d.	n.d.	n.d.	n.d.	n.d.	Yellow solution with brown precipitation
1.7	Clear colourless solution	Clear colourless solution	Clear colourless solution	Clear colourless solution	Deep red solution with precipitation	n.d.
3.5	Clear slightly yellow solution	Clear slightly yellow solution	Clear slightly yellow solution	Clear colourless solution	n.d.	Brown yellow solution with heavy purple precipitation

n.d. = not determined

EXAMPLE 2:**Solubility of omeprazole and omeprazole sodium in polyethylene glycol**

- 5 The solubility of omeprazole (the non-ionised species) has been determined to approximately 0.1 g/l in water and approximately 100 g/l in dichloromethane at room temperature (22 °C).

This means that the solubility of omeprazole is much better in dichloromethane, a less
10 polar solvent, than in water.

Omeprazole sodium on the other hand is freely soluble in water and only slightly soluble in dichloromethane and this, again, is due to the large differences in polarity of the different solvents. Omeprazole sodium is ionised and is therefore much more soluble in water compared with omeprazole and the opposite is applicable in the less polar solvent
15 dichloromethane.

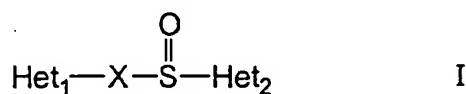
The solubility of omeprazole (the non-ionised species) has been determined to approximately 20 g/l in polyethylene glycol 400. This means that the solubility is much better in polyethylene glycol, a less polar solvent, than in water.

20

Omeprazole sodium is freely soluble in polyethylene glycol 400, which is unexpected and contradictory to what was expected. This means that the solubility in polyethylene glycol is much higher for omeprazole sodium than for omeprazole. The high solubility of omeprazole sodium in polyethylene glycol is very favourable regarding the formulation
25 aspects of a parenteral pharmaceutical product.

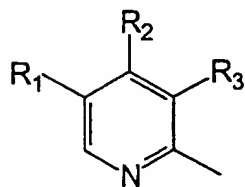
Claims

1. A water free or almost water free, stable liquid formulation comprising polyethylene glycol and a sodium or potassium salt of a H^+ , K^+ -ATPase inhibitor, and optionally pharmaceutically acceptable excipients that are soluble in the formulation.
2. A stable liquid formulation according to claim 1, wherein the sodium or potassium salt of the H^+ , K^+ -ATPase inhibitor is a sodium or potassium salt of a compound with the general formula I, or a sodium or potassium salt of the single enantiomer thereof

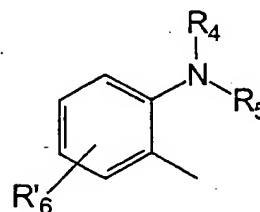


wherein

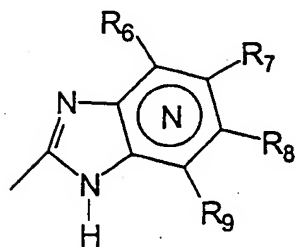
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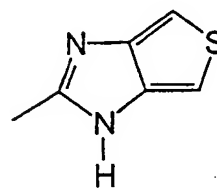
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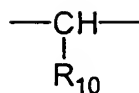
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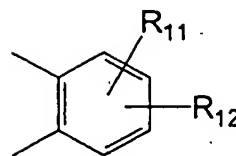
or



X =



or



wherein

N in the benzimidazole moiety means that one of the ring carbon atoms substituted by R₆-

5 R₉ optionally may be exchanged for a nitrogen atom without any substituents;

R₁, R₂ and R₃ are the same or different and selected from hydrogen, alkyl, alkoxy optionally substituted by fluorine, alkylthio, alkoxyalkoxy, dialkylamino, piperidino, morpholino, halogen, phenyl and phenylalkoxy;

10

R₄ and R₅ are the same or different and selected from hydrogen, alkyl and aralkyl;

R₆' is hydrogen, halogen, trifluoromethyl, alkyl and alkoxy;

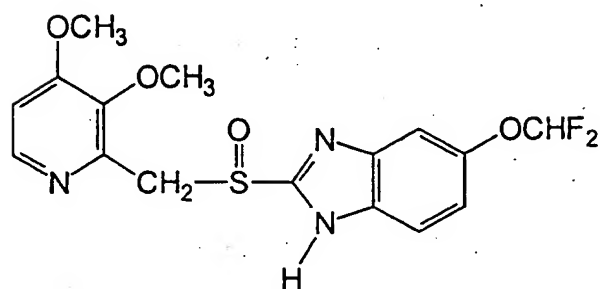
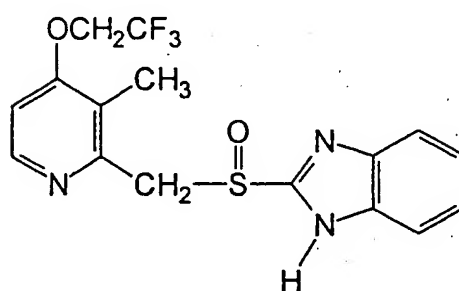
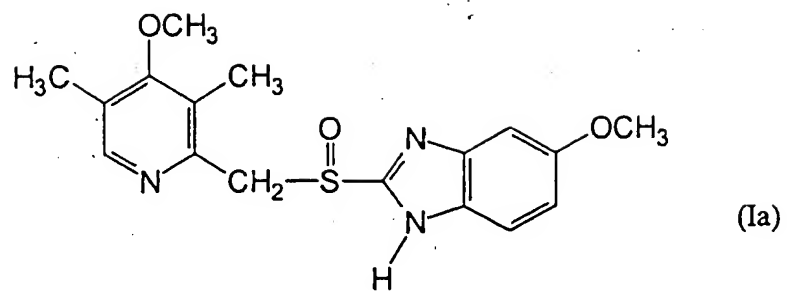
R₆-R₉ are the same or different and selected from hydrogen, alkyl, alkoxy, halogen, halo-

15 alkoxy, alkylcarbonyl, alkoxy carbonyl, oxazolyl, trifluoroalkyl, or adjacent groups R₆-R₉ form ring structures which may be further substituted;

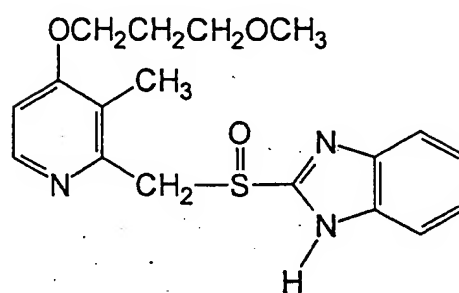
R₁₀ is hydrogen or forms an alkylene chain together with R₃ and

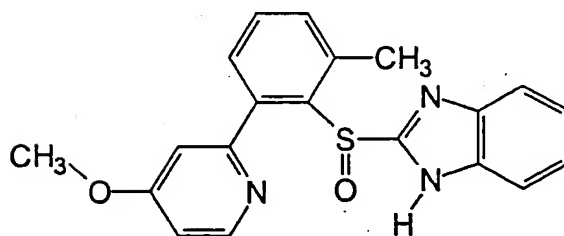
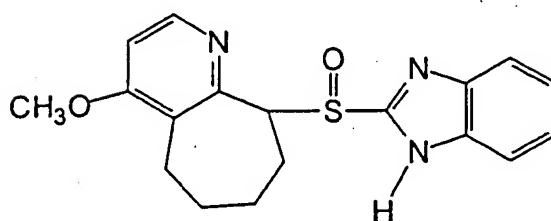
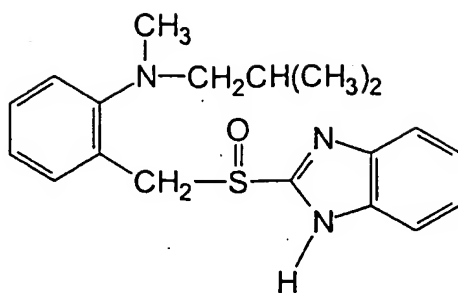
20 R₁₁ and R₁₂ are the same or different and selected from hydrogen, halogen or alkyl.

3. A stable liquid formulation according to any of claims 1 - 2, wherein the sodium or potassium salt of the H⁺, K⁺-ATPase inhibitor is the sodium or potassium salt of a compound with any of the following formulas, or the sodium or potassium salt of the
25 single enantiomer thereof

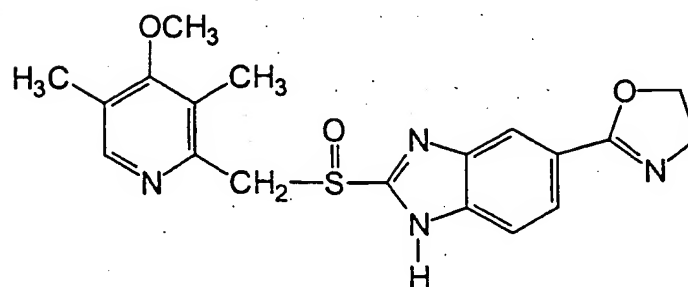
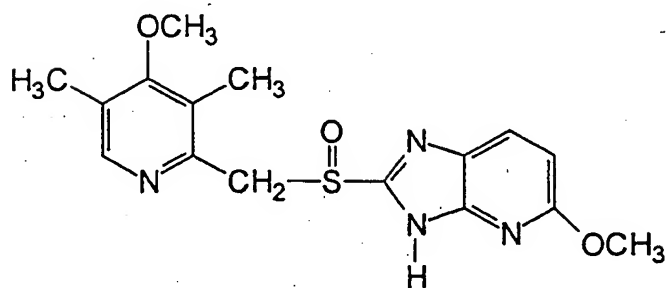


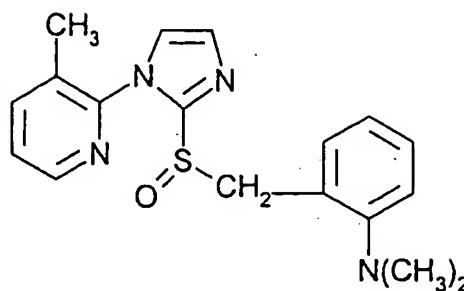
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4. A stable liquid formulation according to any of claims 1 - 3, wherein the compound is a sodium salt.
5. A stable liquid formulation according to any of claims 1-3, wherein the compound is a potassium salt.
6. A stable liquid formulation according to any of claims 1-4 wherein the compound is selected from sodium salt of (*S*)-omeprazole and sodium salt of omeprazole.
7. A stable liquid formulation according to any of claims 1 - 6, wherein the polyethylene glycol is selected from polyethylene glycol 200, 300 and 400.
8. A stable liquid formulation according to claim 7, wherein the polyethylene glycol is polyethylene glycol 400.
9. A stable liquid formulation according to any of claims 1 - 8, wherein the water content of the solution is less than 6% by weight, and preferably less than 3% by weight.
10. A stable liquid formulation according to claim 9, wherein the water content is less than 2 % by weight.
11. A process for the preparation of a stable liquid formulation comprising polyethylene glycol and a sodium or potassium salt of a H^+ , K^+ -ATPase inhibitor, and optionally pharmaceutically acceptable excipients, wherein the sodium or potassium salt of the H^+ , K^+ -ATPase inhibitor and polyethylene glycol are mixed, and optionally excipients that are soluble are added, whereafter the solution optionally is purged with nitrogen to obtain a water content of a pre-set value.

12. A process for the preparation of a stable liquid formulation comprising polyethylene glycol and a sodium or potassium salt of a H^+ , K^+ -ATPase inhibitor, and optionally pharmaceutically acceptable excipients, wherein the H^+ , K^+ -ATPase inhibitor, sodium or potassium hydroxide and polyethylene glycol are mixed, and optionally excipients that are soluble are added, whereafter the solution optionally is purged with nitrogen to obtain a water content of a pre-set value.
13. A process according to claim 10 or 11, characterized in that the process comprises the additional step that the liquid formulation is sterile filtered.
14. The use of a stable liquid formulation according to claim 1 in medicine.
- 10 15. The use of polyethylene glycol and sodium or potassium salt of a H^+ , K^+ -ATPase inhibitor in the manufacture of a stable liquid formulation according to any of claims 1 – 10 to be used in the treatment of gastro intestinal diseases.
16. A method for treating gastrointestinal diseases wherein a stable liquid formulation according to any of claims 1 – 10 is administered to a subject in the need thereof.

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: A61K 31/44, A61K 47/30

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0356143 A1 (TAKEDA CHEMICAL INDUSTRIES, LTD.), 28 February 1990 (28.02.90) --	1-15
A	EP 0444625 A1 (HANMI PHARM. IND. CO., LTD.), 4 Sept 1991 (04.09.91) -- -----	1-15

☐ Further documents are listed in the continuation of Box C.☒ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

12 March 2001

Date of mailing of the international search report

23 March 2001

Name and mailing address of the ISA:

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Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 16
because they relate to subject matter not required to be searched by this Authority, namely:
see next sheet
2. ☒ Claims Nos.: 1
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
see next sheet
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

Box I.1

Claim 16 relate to a method of treatment of the human or animal body by surgery or by therapy/a diagnostic method practised on the human or animal body/Rule. 39.1.(iv). Nevertheless, a search has been executed for this claim. The search has been based on the alleged effects of the compound/composition.

Box I.2

Present claim 1 relates to a compound defined by reference to a desirable characteristic namely H^+ , K^+ - ATPase inhibitors. The claim covers all compounds having this characteristic. The claim also lacks clarity (Article 6 PCT) since the scope of protection covers an undefined amount of compounds with undefined structure. This lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible. Consequently, the search has been carried out for the compounds defined in claim 2.

INTERNATIONAL SEARCH REPORT
Information on patent family members

25/02/01

International Application No.

PCT/SE 00/01992

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